

**USE OF POLYACRYLAMIDE GEL FOR FORMING A CAPSULE
IN THE TISSUE OF THE ORGANISM OF A MAMMAL, A METHOD
OF CULTIVATING CELLS, AND A METHOD OF TREATING
ONCOLOGICAL DISEASES AND DIABETES MELLITUS**

5 The present invention relates to medicine, particularly to immunology and immunooncology, and also to the treatment of diabetes mellitus, predominantly of insulin-dependent diabetes mellitus. More particularly, the invention relates to the problem of vaccination against tumor cells and of vaccinothrapy of oncological diseases and to a new method of treating diabetes mellitus. Furthermore, the invention proposes a special
10 capsule for use in methods of treating, owing to which the treatment proves to be substantially more effective.

It is known that the problem of transplanting organs, tissues, cell cultures of mammals involves difficulties associated with the possibility of xenogenous tissue-and-cell agents to "take root" in the recipients' organism. The existing methods of transplanting
15 allo-, hetero- and xenotransplants require either powerful immunosuppressive therapy of the recipient or original procedures. These latter comprise methods of transplanting cells of various organs of human and animal fetuses, i.e., the effect of undeveloped species specificity is employed. In such a manner, e.g., islet cell cultures of the pancreas of 24—26 weeks' human fetuses are transplanted into the parenchyma of the liver or into the portal
20 vein in experiments with rats.

The place of introducing cells may be splenic pulp or muscles of the anterior abdominal wall. Cases of treating humans suffering from diabetes mellitus by a similar method are known (Skaletskii N.N., "The Effect of Cultivating Islet Cells of the Pancreas on Their Survival in the Organism of a Xenogenous Recipient", in: All-Russian
25 Conference on Transplantation of Organs, 1995, pp. 219—220 (in Russian)).

The transplantation of Leydig's cells into testicular tissue to males for treating infertility is of interest, because the rejection reaction does not occur due to the presence of a hematotesticular barrier (Zybin D.V., "Method of Treating Patients with Dysfunction of Male Sexual Sphere by Transplantation Techniques", RF Patent 2026643 of 20.01.95).

30 As a result of both above-described methods, good results were obtained in preserving the viability and activity of transplanted cells. However, the first method makes it possible to use only embryonal cells, this, for obvious reasons, involving a number of difficulties; the second method of cell therapy proves to be applicable only to male individuals.

There is known a method of vaccination and vaccinothrapy of tumors with the help of live cells. The employed cells are hybridomas or transformed cells, allogenic or autogenic. A disadvantage of such cells is their short-time existence in the organism and, correspondingly, a low immunizing effect (B.E. Souberbielle, M. Westby, S. Ganz, and J.

5 Kayaga, Comparison of four strategies for tumor vaccination in the B-16

F 10 melanoma model. Gene Therapy 1998, 1447—1454).

Known in the art is a method of transplantation of pancreatic islets (PIs) using microencapsulation.

10 The method consists in introducing (allo- or xenogenic) PIs encapsulated in spheres of an alginate gel. The spheres are implanted intraperitoneally. The implantation of spheres completely replaces insulin therapy during 175 days, but in this case immunosuppressive therapy is concurrently used. Bovine PIs are administered to rats with induced diabetes without immunosuppression. Normoglycemia is maintained from several weeks to one month.

15 A certain inconvenience of the method is the necessity of using immunosuppressive therapy. Definite difficulties are involved in preparing capsules in vitro (Lanza R.P., Esker D.M., and Marsh J.P., Transplantation of islets using microencapsulation studies in diabetic rodents and dogs. J. Mol. Med., 1999 Jan. 77(1): 206—10).

20 Also known is a method of vaccination and vaccinothrapy of tumors with the help of live cells. It consists in producing hybridomas of tumor cells and allogenic dendritic cells (or macrophages). The obtained hybridomas are used as vaccine preparations.

25 However, this method also features a low immunizing effect, caused by the short-time existence of the introduced cells in the recipient's organism (Gajewsky T.F. and Fallarino F., Rational development of tumor antigen-specific immunization in melanoma. Therapeutic Immunology, 1997, 2, 211—225).

Therefore, the problem of increasing the life-span of transplanted cells and, as a consequence, of enhancing the immunizing effect, as well as obviating immunosuppressive therapy is now as before topical in this field of the art.

30 Specialists are aware that the problem of treating diabetes mellitus is also closely connected with the positive solution of the question of transplanting cells, whose successful solution will in many respects make for the desired effectiveness of the method of treating.

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For instance, a method of treating diabetes mellitus is known, according to which implantation of cells of benignant human insulinoma is carried out, the material containing pancreatic β -cells is implanted into the musculus rectus abdominis (RF Patent 2004247).

However, problems which arise in connection with combating the predominance of the growth of fibroblasts when using a culture of β -cells for transplantation, and the necessity of precise control of the insulin production by a particular fraction of the insulinoma cell culture, called for due to the fact that tumor cells whose functional activity may vary substantially are used as the implant, offer a hindrance to the wide use of the method.

Also known is a method of treating diabetes mellitus by transplanting material containing pancreatic β -cells (RF Patent 2135193). The method of treating diabetes mellitus, predominantly insulin-dependent one, is carried out using material containing pancreatic β -cells of mammals, produced with the use of the β -cells migration phenomenon.

The material containing pancreatic β -cells is transplanted into different organs and tissues; intramuscularly, into the musculus rectus abdominis, into the liver (into the parenchyma or via the portal vein), into the splenic pulp, into the spleen artery, into the abdominal cavity, into the greater omentum, into a specially created muscular pocket.

The short period of time during which β -cells produce an insulin donor in the recipient's organism because of the effect of rejection of heterogeneous cells necessitates considerable immunosuppressive therapy.

Figure 1 illustrates the dynamics of testosterone in the blood serum of Vistar-line rats to which Leydig's cells of young pigs (rows 1 and 2) and of green monkeys (rows 3 and 4) are implanted. Rows 1 and 3 show control results (cells are introduced subcutaneously, rows 2 and 4 show experimental results (introducing cells into a formed polyacrylamide capsule).

The present invention is directed to overcoming the above-indicated problems. The authors of the present invention unexpectedly discovered that long-time maintaining the viability of transplanted cells, including heterogeneous ones, in the recipient's organism can be provided by using a polyacrylamide gel capsule being formed in vivo in the organism of a mammal (including a human) in need of therapy with such cells.

So, one of the aspects of the present invention is the use of a polyacrylamide gel for preparing in the organism of a mammal a polyacrylamide capsule formed therein in vivo, which capsule can later be used for cultivating cells transplanted thereinto.

Unexpectedly to the authors of the present invention, cells injected into the above-
5 said capsule proved to be capable of preserving viability for a long period of time (to 100 days and more) and of producing compounds necessary for the treatment.

Another aspect of the invention is, therefore, a method of cultivating cells necessary for the treatment in the organism of a patient in need of such treatment. The cultivation of cells is preceded by the injection of a polyacrylamide gel into the organism
10 of a mammal; by the formation of a gel capsule during a definite period of time in the organism of a mammal; and by the injection into said capsule of a required amount of cells to be transplanted.

Long-time survival of the cells in the cultivation thereof inside the patient's body proves to be useful for the treatment of a number of diseases, which requires
15 transplantation of autologous or heterologous cells producing biologically active compounds whose deficiency in the organism aggravates or initiates the disease.

The third aspect of the present invention is a method of treating diseases for which immunization with an antigen is indicated, this antigen under usual conditions requiring intensive immunosuppressive therapy.

20 The next aspect of the invention is a method of treating diabetes mellitus, predominantly insulin-dependent one, consisting in that an effective amount of pancreatic β -cells is introduced into a polyacrylamide gel capsule preformed in the organism of a patient.

One more aspect of the present invention is a method of cultivation and
25 modification of heterogeneous cells (tumor cells, Leydig's cells, etc.) in the organism of a mammal with a view to their subsequent use for producing a vaccine preparation. The definition "modification of heterogeneous cells" should be understood as lowering the proliferative activity and immunizing action on the organism.

The invention will further be disclosed in detail by examples of its preferred
30 embodiment, these examples being given by way of illustration only and should not be used for limiting the scope of claims. A person skilled in the art may find a considerable number of possibilities for complementing or modifying the invention which will preserve the above-indicated advantages and will be encompassed by the set of claims.

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In a general form the invention is carried out as follows.

A connective-tissue capsule according to the present invention may be formed, for instance, by way of subcutaneous injection of a polyacrylamide gel (PAAG) (in a volume of 1.0—5 ml) to animals, e.g., to Vistar-line rats (in a volume of 1.0—3 ml) or (in a volume of 0.5—1 ml) to mice of C57BLACK and BALB/C lines or to a mammal such as a human (in a volume of 1.0—3.0 ml). Individuals of different sexes may participate in the experiment. Leydig's cells of pubescent young pigs, rats, and green monkeys or tumor cells can be introduced into the gel capsule. Animals to which cells are injected subcutaneously serve as control.

A suspension of viable Leydig's cells from the testicles of pubescent young pigs, rats, and green monkeys is prepared by using solutions containing a nutrient substrate for cells, in particular, compositions of standard Eagle's medium, medium 199, Hanks' solution, etc.

The essence of the proposed method of treating patients suffering from diabetes mellitus is the long-time existence of and production of insulin by pancreatic β -cells of the donor in the recipient's organism, this being achieved by preliminary subcutaneous injection of a polyacrylamide gel to the patient and subsequent transplantation of β -cells into the formed capsule.

The material for the transplantation of β -cells is obtained from the pancreas of mammals (newborn pigs, rabbits, pubescent green monkeys). Cultivation is carried out using standard media and solutions. For preserving β -cells in the active state, a method of sparing enzymatic treatment of the pancreas is employed, which consists in alternating contacts of the tissue with the enzyme and the nutrient medium. As a result of the treatment steps, fragments of the pancreas tissue and β -cells are introduced into cultivation vessels without centrifugation. Disaggregation of the tissue is carried out with a 0.1—0.25% solution of trypsin and chenopsin in different sequences, depending on the donor material. The enzymatic treatment of the tissue is completed during its contacts with the medium, using a "Biotech-m" flask, which provides for controlled stirring of the suspension on a magnetic table.

The resulting cell material is injected to the patient into the connective-tissue capsule which is formed by the polyacrylamide gel preliminarily administered subcutaneously. The amount of the cells depends on the gravity of the recipient's disease.

The Examples which follow illustrate carrying the invention into effect

Example 1

A culture of Leydig's cells of newborn pigs in the volume of 0.5 ml with the concentration of cells of 5 million per ml is injected into a polyacrylamide gel capsule to females of Vistar-line rats.

To the control group of animals of the same line, sex and anthropological data a culture of Leydig's cells of newborn pigs is administered subcutaneously. Before injecting the cell culture, the content of testosterone in the blood serum of the animals is measured. Subsequent measurements of testosterone in the blood serum are carried out with different intervals during 7 months in the control and experimental animals simultaneously.

The number of animals in control and in the experiment was 2 individuals in each. Fig. 1 (rows 1 and 2, respectively).

Example 2

A culture of Leydig's cells of pubescent green monkeys in the volume of 0.5 ml with the concentration of cells of 5 million per ml is injected into a polyacrylamide gel capsule to females of Vistar-line rats.

To the control group of animals of the same line, sex and anthropological data a culture of Leydig's cells of pubescent green monkeys is administered subcutaneously. Before injecting the cell culture, the content of testosterone in the blood serum of the animals is measured. Subsequent measurements of testosterone in the blood serum are carried out with different intervals during 7 months in the control and experimental animals simultaneously.

The number of animals in control and in the experiment was 2 individuals in each. Fig. 1 (rows 1 and 2, respectively).

After 7 months of observations the animals are sacrificed and a histological investigation is carried out. It indicates the presence of a large amount of viable Leydig's cells, so that a conclusion can be drawn about the possibility of vital activity of xeno- and heterogeneous cells in the recipient's organism with the use of a gel.

Example 3

PAAG in the volume of 0.5 ml is administered subcutaneously to an experimental group of mice of the BALB/C line (6 individuals). Tumor cells of murine melanoma B-16 are injected into the gel in the volume of 1 ml with the concentration of cells of 1 million cells per ml. Cells of murine melanoma B-16 in the volume of 1 ml with the concentration

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of cells of 1 million per ml are administered subcutaneously to a control group of mice of the BALB/C line (6 individuals).

It is known that in mice of the BALB/C line the tumor of murine melanoma B-16 does not grow. In the control group of the animals the tumor growth was found in none of the 6 individuals. In the experimental group of the animals, by way of palpatory examination, a growth of tumor in the PAAG was noted in all the 6 individuals. By the 60th day the experimental animals with the murine melanoma B-16 in the gel are sacrificed. The gel with tumor cells is extracted under aseptic conditions and transferred to a monolayer culture on the nutrient medium RPMI-1640 with 10% fetal serum. Fragments of the capsule with the tumor cells are fixed in a neutral solution of formalin, and a histological investigation is carried out, which allows one to judge about a higher differentiation of melanoma cells and the loss of the proliferative activity by them (Table 1 [1—2]).

TABLE 1. COMPARISON OF THE GROWTH OF MELANOMAS B-16 (MURINE) AND SKMEL 28 (HUMAN) IN MICE OF BALB/C AND C57BLACK LINES

LINE OF MICE	TUMOR STRAIN	GROWTH OF MELANOMA	SPAN OF LIFE	META-STASES
BALB/C + GEL	B-16	+	60 DAYS (observa-tion period)*	-
BALB/C	B-16	-	> OBSERVA-TION PERIOD	-
C57BLACK + GEL	SKMEL28	+	> OBSERVA-TION PERIOD	-
C57BLACK	SKMEL28	-	> OBSERVA-TION PERIOD**	-

* Animals with tumors grown in the gel are sacrificed. The tumor cells isolated from them are used in the next experiment (Table 2).

** Mice are used further in the experiment for estimating immunity against melanoma B-16 (Table 3).

Example 4

A culture of cells prepared as in Example 1 is administered subcutaneously in the amount of 1 ml with the concentration of cells of 1 million to mice of the C57BLACK line (6 individuals).

It is known that in mice of the C57BLACK line the tumor of murine melanoma B-16 grows in 100% of cases, death of the animals occurs on the 20—25th day in 100% of cases.

To the control group of C57BLACK mice a culture of cells of murine melanoma B-16 is administered in the amount of 1 ml with the concentration of cells of 1 million.

In experimental mice the appearance of symptoms of tumor growth is noted in 60—65 days; in control mice, in 20—23 days

(Table 2).

TABLE 2. TUMORIGENIC ACTIVITY OF MELANOMA B-16, CULTIVATED IN A GEL CAPSULE, IN MICE OF BALB/C LINE, GRAFTED TO MICE OF C57BLACK LINE

No.	TUMOR STRAIN	TIME OF TUMOR APPEARANCE	LIFE-SPAN OF MICE	PRESENCE OF METASTASES
1	Melanoma from gel of mice of BALB/C line (B-16-X)	30 DAYS	60 DAYS	+
2	B-16 (control)	7 DAYS	22 DAYS	+

Example 5

PAAG in the volume of 0.5 ml is injected subcutaneously to mice of the C57BLACK line (6 individuals). A cell culture of human melanoma SKMEL is injected into the gel in the amount of 1 ml with the concentration of cells of 1 million.

The control group of mice of the same line (6 individuals) is administered subcutaneously a cell culture of human melanoma SKMEL28 in the volume of 1 ml with the concentration of cells of 1 million.

It is known that the cell culture of human melanoma does not grow in mice in 100% of cases. In the experimental group of animals the tumor growth in the gel is determined by palpation on the 15-20th day after the injection. In the control animals no tumor growth is found (Table 1 [3—4]).

Example 6

The group of experimental animals (6 individuals), described in Example 3, is administered subcutaneously a cell culture of murine melanoma B-16 in the volume of 1 ml with the concentration of cells of 1 million. The control group of mice of the C57BLACK line (6 individuals) is administered subcutaneously a cell culture of murine melanoma B-16 in the volume of 1 ml with the concentration of cells of 1 million.

In the control animals subcutaneous melanomas approximately 3—5 cm in diameter develop on the 7—15th day. In the experimental animals no symptoms of tumor are detected during the same period of time (Table 3).

TABLE 3. IMMUNOGENIC ACTIVITY OF HUMAN MELANOMA SKMEL28 FOR MICE

No.	TUMOR STRAIN	TUMOR APPEARANCE TIME	DEATH OF ANIMALS
5	B-16	-	> 60 DAYS
6	B-16	7—15 DAYS	18—20 DAYS

So, the results presented hereinabove give grounds to believe that the method of cultivating heterogeneous cells in PAAG in vivo, whereby the proliferative activity of tumor cells lowers and the cultivated cells produce an immunizing effect on the organism, can be used for vaccination and vaccinotherapy.

Example 7

Female patient F., aged 37. Insulin-dependent diabetes mellitus was diagnosed 11 years ago, one year after the parturition. The pregnancy was aggravated by toxicosis during the second half of the pregnancy period, by nephropathology, a considerable, to 26 kg, gain in weight. The character of the disease has been unstable all these years, and considerable efforts were required in choosing adequate insulin therapy. The use of exogenous insulin was varied from 58 units per day to 30 units per day. During the last two years pathological changes in the kidneys were diagnosed, defined as diabetic nephropathy. In urinalyses a 10—12-fold increase of the upper boundary of proteinuria is noted. The arterial pressure increases to 170/110 mm Hg.

A culture of pancreatic cells of newborn rabbits was administered to the patient subcutaneously, by injecting the culture into a preliminarily formed capsule. Already 7 days later the patient noted an improvement in the general condition, a reduction of the

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Example 8

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secondary complications of diabetes mellitus: rethino-, neuro-, nephropathies. The method makes it possible to improve appreciably the quality of life of patients. As a rule, the therapeutic effect lasts for 10—20 months, depending on the severity of the disease.

5 The amount of cells to be transplanted also depends on the severity of the course of diabetes mellitus, particularly on the quantity of exogenous insulin taken by the patient.

10 The proposed invention makes it possible, using "a polyacrylamide gel", by introducing it into the organism of a mammal, to form in vivo a capsule, which later on, being injected with viable cells for transplantation, functions as a chamber for cultivating producer cells during a long period of time, required for treating with a compound produced by the cell, when said compound, releasing from this artificially formed chamber, produces the desired effect on the organism of a patient. The use of a polyacrylamide gel for the indicated purposes allows maintaining the viability of cultivated cells for a long time and thereby provides a prolonged curative effect. Said use makes it possible to rule out immunosuppressive therapy and may find very extensive application in
15 practical medicine.

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